4376 FIRST AID KIT- 4376 first aid kit Honeywell Safety Products USA, INC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

4376 First Aid Kit (BZK wipe,FABC, neomycin, alcohol wipes, sting relief, ASA, aypanal-Z019807)

Alcohol Wipe Active ingredient

Isopropyl alcohol 70%

Alcohol Wipe Purpose

First aid antiseptic

Alcohol Wipe

Uses

• first aid to help prevent infection in minor cuts, scrapes, and burns

Alcohol Wipe Warnings

For external use only

Do not use

- in the eyes
- over large areas of the body

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burn

When using this product

• do not use longer than one week unless directed by a doctor

Stop use and consult a doctor

• if condition persists or gets worse

Keep out of reach of children

• If swallowed, get medical help or contact a Poison Control Center right away.

Alcohol Wipe Directions

- clean the affected area
- apply wipe to affected area 1 to 3 times daily
- may be covered with a sterile bandage
- discard wipe after single use

Alcohol Wipe Other information

store at room temperature 15 0 to 25 0 C (59 0 to 77 0 F)

Alcohol Wipe Inactive ingredient

water

Alcohol Wipe Questions

1-800-430-5490

Aspirin

Active ingredient (in each tablet)

Aspirin 325 mg (NSAID)* *nonsteroidal anti-inflammatory drug

Aspirin

Purpose

Pain reliever/fever reducer

Aspirin

Uses

temporarily reduces fever and relieves minor aches and pains associated with:

- a cold
- headache
- toothache
- muscular aches
- backache
- minor pain of arthritis
- premenstrual and menstrual periods

Aspirin

Warnings

Reye's syndrome: Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's syndrome, a rare but serious illness.

Allergy alert: Aspirin may cause a severe allergic reaction which may include:

- hives
- facial swelling

- asthma (wheezing)
- shock

Stomach bleeding warning: This product contains an NSAID, which may cause severe stomach bleeding. The chance is higher if you:are:

- age 60 or older
- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
- have 3 or more alcoholic drinks every day while using this product
- take more or for a longer time than directed

Do not use

• if you are allergic to aspirin or any other pain reliever/fever reducer

Ask a doctor before use if

- stomach bleeding warning applies to you
- you have a history of stomach problems such as heartburn
- you have high blood pressure, heart disease, liver cirrhosis or kidney disease
- you are taking a diuretic
- you have asthma

Ask a doctor or pharmacist before use if you are

• taking a prescription drug for diabetes, gout or arthritis

Stop use and ask a doctor if

- you experience any of the following signs of stomach bleeding:
- feel faint
- vomit blood
- have bloody or black stools
- have stomach pain that does not get better
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present in the painful area
- ringing in the ears or loss of hearing occurs
- any new symptoms appear

If pregnant or breast-feeding,

If pregnant or breat-feeding, ask a health professional before use. It is especially important not to use aspirin during the last three months of pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

Keep out of reach of children.

• In case of overdose, get medical help or contact Poison Control Center right away.

Aspirin

Directions

- drink a full glass of water with each dose
- adults and children 12 years of age and older: take 1 or 2 tablets every 4 hours while symptoms last,

not more than 12 tablets in 24 hours

• children under 12 years of age: consult a doctor

Aspirin

Other information

- store at room temperature 15° 30°C (59° 86°F)
- TAMPER EVIDIENT PACKETS
- DO NOT USE IF OPEN OR TORN

Aspirin

Inactive ingredients

corn starch, croscarmellose sodium*, hypromellose*, microcrystalline cellulose*, mineral oil*, polyethylene glycol*, povidone, propylene glycol, silicon dioxide, stearic acid*, titanium dioxide* *may contain these ingredients

Aspirin

Questions or Comments

1-800-430-5490

Sting Re;ief Active ingredient

Ethyl alcohol 50.0%

Lidocaine HCl 2.0%

Sting Relief Purpose

Antiseptic

Topical pain relief

Sting Relief

Uses

• prevent infection in minor scrapes, and temporary relief of itching of insect bites

Sting Relief

Warnings

For external use only

Flammable, keep away from open fire or flame

Do not use

- over large areas of the body
- in eyes
- over raw or blistered areas

Stop use and ask a doctor

• if conditions worsen or persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

sting Relief

Directions

- adults and children 2 years and older: Apply to cleaned affected area not more than 3 times daily.
- children under 2 years of age: consult a doctor.

Sting Relief

Inactive ingredients

benzalkonium chloride, menthol, and purified water

Sting Relief

Questions or Comments?

1-800-430-5490

BZK

Active ingredient

Benzalkonium chloride 0.13% w/v

BZK

Purpose

First aid antiseptic

BZK

Uses

Antiseptic cleansing of face, hands, and body without soap and water

BZK

Warnings

For external use only

Do not use

- in the eyes or over large areas of the body
- on mucous membranes
- on irritated skin
- in case of deep puncture wounds, animal bites or serious burns, consult a doctor
- longer than 1 week unless directed by a doctor

Stop use and ask a doctor if

- if irritation, redness or other symptoms develop
- the condition persists or gets worse

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

BZK

Directions

• tear open packet and use as a washcloth

BZK

Other information

- store at room temperature 15 0 to 30 0 C (59 0 86 0 F)
- do not reuse towelette

BZK

Inactive ingredient

water

BZK

Questions

1-800-430-5490

Neomycin

Active ingredient (each gram contains)

Neomycin sulfate (5 mg equivalent to 3.5 mg Neomycin base)

Neomycin

Purpose

First aid antibiotic

Neomycin

Uses

- first aid to help prevent infection in
- minor cuts
- scrapes
- burns

Do not use

- in the eyes
- over large areas of the body

Neomycin

Warnings

For external use only

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- the condition persists or gets worse
- a rash or other allergic reaction develops
- you need to use longer than 1 week

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

Neomycin

Directions

- clean the affected area
- apply a small amount of the product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily
- may be covered with a sterile bandage

Neomycin

Other information

• store at 15 0 to 25 0 C (59 0 to 77 0 F)

Neomycin

Inactive ingredient

petrolatum

Neomycin *Questions*

1-800-430-5490

Aypanal EX

Active ingredient

Acetaminophen 500 mg

Aypanal EX

PUrpose

Pain reliever/fever reducer

Aypanal EX

Uses

• temporarily relieves minor aches and pains due to the common cold and headache - temporarily reduces fever

Aypanal EX

Warnings

Liver Warning: This product contains acetaminophen. Severe liver damage may occur if you take:

• more than 4,000 mg in 24 hours, which is the maximum daily amount.

- with other drugs containing acetaminophen
- adult has 3 or more alcoholic drinks every day while using this product

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If skin reaction occurs, stop use and seek medical help right away

Do Not Use

• with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

Ask a doctor before use if you have

liver disease

Ask a doctor or pharmacist before use if

• you are taking the blood thinning drug warfarin

Stop use and ask a doctor if

- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- new symptoms occur
- redness or swelling is present
- If pregnant or breastfeeding,
- ask a health professional before use.

Keep out of reach of children.

Overdose warning: In case of loverdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Avpanal EX

Directions

- do not take more than directed (see overdose warning)
- adults and children 12 years of age and over: Take 2 tablets with water every 6 hours while symptoms last.
- do not take any more than 8 tablets in 24 hours.
- children under 12: consult a doctor

Aypanal EX

Other information

- store at room temperature 15 0 -30 0 C (59 0 -86 0 F)
- TAMPER EVIDENT- DO NOT USE IF OPEN OR TORN

Aypanal EX

Inactive ingredients

microcrystalline cellulose, povidone, sodium starch glycolate, starch, stearic acid

Aypanal EX

Questions or Comments?

1-800-430-5490

FABC

Active ingredient

Benzalkonium chloride 0.13%

Lidocaine HCl 0.5%

FABC

Purpose

First Aid antiseptic

External analgesic

FABC

Uses

- prevent skin infection
- for temporary relief of pain associated with minor burns

FABC

Warnings

For external use only

Do not use

- in or near the eyes
- if you are allergic to any of the ingredients
- lin large areas of the body, particularly over raw surfaces or blistered areas
- for more than 10 days

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days or clear up and occurs again within a few days

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

FABC

Direcctions

- adults and children 2 years of age and older:
- clean the affected area
- apply a small amount of this product (equal to the surface area of the tip of a finger) onto affected

- area 1 to 3 times daily
- may be covered with a sterile bandage
- children under 2 years of age: consult a doctor

FABC

Other information

- tamper evident sealed packets
- do not use if packet is opened or torn

FABC

Inactive ingredients

aloe barbadensis juice, cetyl alcohol, diazolidinyl urea, edetate disodium, glycerin, glyceryl stearate SE, methylparaben, mineral oil, PEG-100, propylene glycol, propylparaben, stearic acid, trolamine, water

FABC

Questions

1-800-430-5490

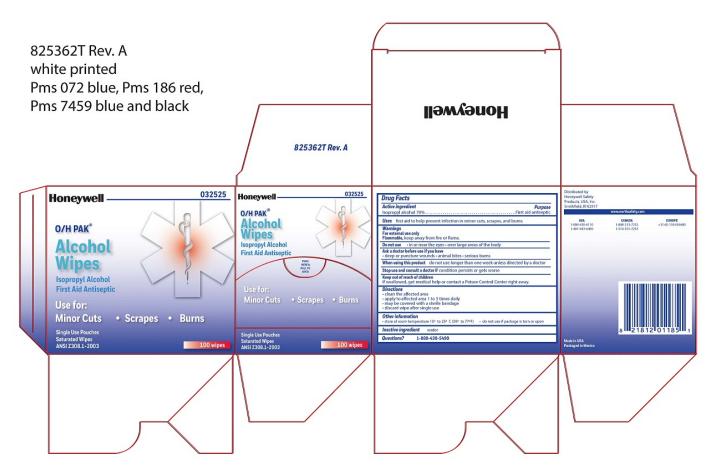
4376

Z019807 KIT CONTENTS

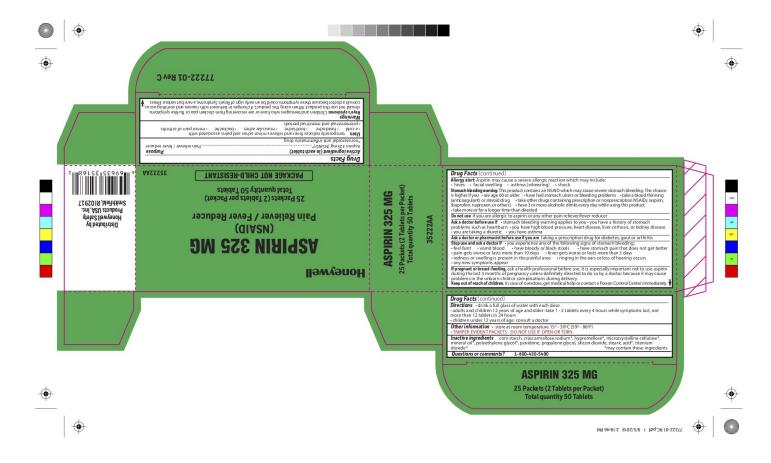
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- 1 FIRST AID GUIDE ASHI
- 1 EMERGENCY SURVIVAL BLANKET
- 1 GAUZE CLEAN-WRAP BDGE N/S 2"
- 10 CTA 6" SINGLE TIP 100/BAG
- 1 SCISSOR BDGE 4" RED PLS HDL
- 1 FANNY PACK RED FAK LOGO EMPTY
- 4 TONGUE BLADES SR WRAPPED 250
- LBL STOCK 6-3/8"X4"
- 1 LBL STOCK 3"x1-7/8"
- 1 LBL CONTS 6 3/4"X3 1/2" ID B
- 6 BZK ANTISEPTIC WIPE, BULK
- 1 PR LRG NITRILE GLVES ZIP BAG
- 3 FIRST AID BURN CREAM 1.0GR PKT EACH
- 3 POUCH NEOMYCIN ANTIBIOTIC .9 G
- 4 ADH BANDAGE BUTTERFLY 1980000
- 6 WIPE ALCOHOL PREP IPA 70% (DUKAL)
- 2 LANCET BLADE #11 STERILE
- 10 ADH BANDAGE 3/8" X 1 1/2" DNX
- 3 SAFETEC STING RELIEF WIPES BULK

- 1 COLD PACK UNIT 4"X6" BULK
- 4 GAUZE PADS 2"X2" 12PLY
- 2 GAUZE PADS 4"X4" 12PLY
- 15 PLASTIC BANDAGE 3/4" X 3"
- 1 WOVEN KNUCKLE BANDAGE
- 1 ZIP-LOCK BAG 5" X 5" .002
- 2 AYPANAL BULK 2/PK
- 2 ASPIRIN BULK 2/PK

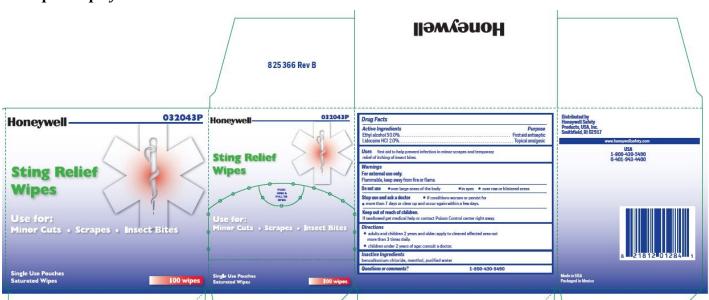
Alcohol Wipe Label



As pirin Principal Display Panel

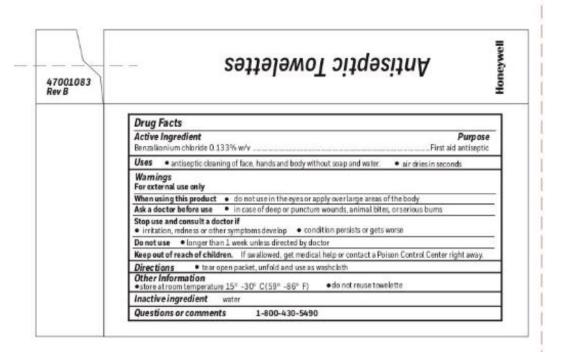


Sting Relief Principal Display Panel



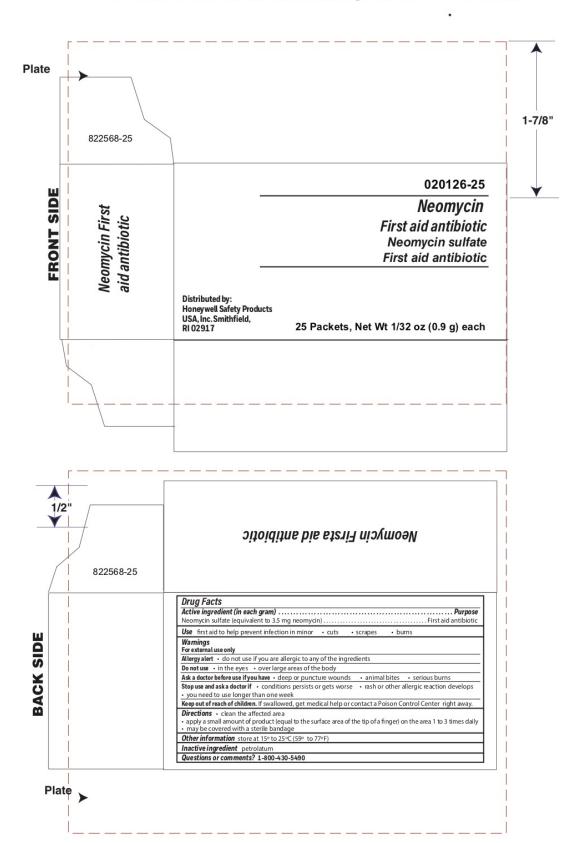
BZK Principal Display Panel

S	Honeywell	
lette		02-16-35MD
оме		Antiseptic Towelettes
Antiseptic Towelettes		Benzalkonium chloride First aid antiseptic
tise		Six-Saturated Towelettes
An	Distributed by Honeywell Safety Products USA, Inc. Smithfield, RI 02917	



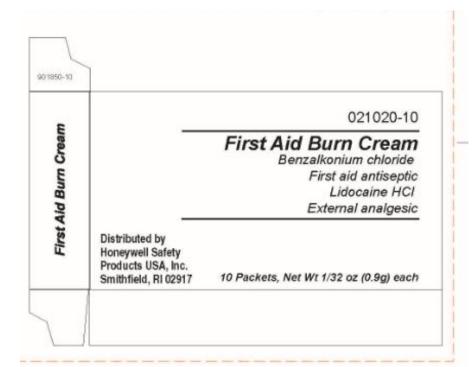
Neomycin Principal Display Panel

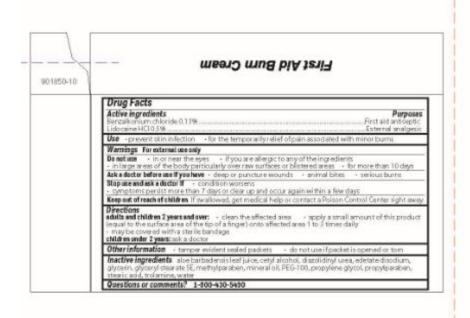
796041-25 Rev A Unit Carton Printing Plate for "C" size carton.



Principal Display Panel -----**EXTRA STRENGTH** Honeywell Honeywell Honeywell Honeywell Drug Facts Active ingredient (in each lable) Active in a discovery in a contract of the active in a contract (in each lable) Active in a discovery in a contract (in each lable) Active in a Distributed by Honeywell Safety Products USA, Inc. Smithfield, RI 02917 EXTRA STRENGTH Drug Facts (continued) Other information -store at room temperature 15:-30" C 150" Pers Acetaminophen 500mg Pain Reliever - Fever Reducer A Complete Line of Unit Dose Tablet Packs Aypanal Extra Strength ain Reliever - Fever Reducer cetaminophen 500 mg

FABC Principal Display Panel





4376 Kit Label Z019807



Distributed by Honeywell Safety Products USA, Inc. Smithfield, RI 02917

APPROVED

4376 FIRST AID KIT

4376 first aid kit kit

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0498-4376

Packaging

ı	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:0498-4376-01	1 in 1 KIT: Type 0: Not a Combination Product	10/18/2018	

Quan	Quantity of Parts		
Part #	Package Quantity	Total Product Quantity	
Part 1	3 PACKET	2.7 g	
Part 2	6 POUCH	2.4 mL	
Part 3	2 PACKET	4	
Part 4	3 PACKET	2.7 g	
Part 5	6 PACKET	8.4 mL	
Part 6	3 POUCH	1.2 mL	
Part 7	2 PACKET	4	

Part 1 of 7

FIRST AID BURN

benzalkonium chloride, lidocaine hydrochloride cream

Product Information

Item Code (Source) NDC:0498-0903

Route of Administration TOPICAL

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g			
LIDO CAINE HYDRO CHLO RIDE (UNII: V13007Z41A) (LIDO CAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g			

Inactive Ingredients		
Ingredient Name	Strength	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
CETYL ALCOHOL (UNII: 936JST6JCN)		
PEG-100 STEARATE (UNII: YD01N1999R)		
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)		
TROLAMINE (UNII: 9O3K93S3TK)		
WATER (UNII: 059QF0KO0R)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
METHYLPARABEN (UNII: A2I8 C7HI9 T)		
GLYCERYL MONOSTEARATE (UNII: 230 O U9 XXE4)		
LIGHT MINERAL OIL (UNII: N6K5787QVP)		
GLYCERIN (UNII: PDC6A3C0OX)		
PROPYLPARABEN (UNII: Z8IX2SC1OH)		

Packaging				
l	# Item Cod	e Package Description	Marketing Start Date	Marketing End Date
l	1	0.9 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Infor	mation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/20/2017	

Part 2 of 7

ALCOHOL WIPE

isopropyl alcohol swab

Product Information

Item Code (Source) NDC:0498-0143

Route of Administration TOPICAL

Active Ingredient/Active Moiety

ı	retive ingredient/retive withery			
I	Ingredient Name	Basis of Strength	Strength	
I	ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.7 mL in 1 mL	

Inactive Ingredients

Ingredient Name Strength

WATER (UNII: 059QF0KO0R)

Packaging

ı		0 0			
l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	_				

1 NDC:0498-0143-04 0.4 mL in 1 POUCH; Type 0: Not a Combination Product

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/18/2018	

Part 3 of 7

ASPIRIN

aspirin tablet

Product Information

Item Code (Source)	NDC:0498-0114
Pouts of Administration	ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)	ASPIRIN	325 mg	

Inactive Ingredients		
Ingredient Name	Strength	
CELLULOSE, MICRO CRYSTALLINE (UNII: OP1R32D61U)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
STARCH, CORN (UNII: O8232NY3SJ)		
PO VIDO NE (UNII: FZ989 GH94E)		
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)		
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)		
HYPROMELLOSE 2208 (100 MPA.S) (UNII: B1QE5P712K)		
MINERAL OIL (UNII: T5L8T28FGP)		
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		

Product Characteristics				
Color	white	Score	2 pieces	
Shape	ROUND	Size	10 mm	
Flavor		Imprint Code	FR21	
Contains				

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0114-01	2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part343	09/18/2018		

Part 4 of 7

NEOMYCIN

antibiotic ointment

NDC:0498-0730
TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297)	NEOMYCIN SULFATE	3.5 mg in 1 g	

Inactive Ingredients	
Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

ı	P	ackaging			
l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:0498-0730-01	0.9 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	03/31/2010	

Part 5 of 7

ANTISEPTIC TOWELETTE

benzalkonium chloride liquid

Product Information	
Item Code (Source)	NDC:0498-0501
Route of Administration	TOPICAL

Active Ingredient/Active Moiety			
Ingre	dient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5 UNII:7N6JUD5X6Y)	UM2KM3W7) (BENZALKONIUM -	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

l	Packaging					
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1 NDC:0498-0501-00	1.4 mL in 1 PACKET; Type 0: Not a Combination Product				

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/22/2017	

Part 6 of 7

STING RELIEF PAD

ethyl alcohol, lidocaine swab

Product Information

Item Code (Source)NDC:0498-0733

Route of Administration TOPICAL

Active Ingredient/Active Moiety

ricare ingredient retire in oreign				
Ingredient Name	Basis of Strength	Strength		
LIDO CAINE HYDRO CHLO RIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98 PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.5 mL in 1 mL		

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
MENTHOL (UNII: L7T10 EIP3A)	
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)	

Packaging

l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:0498-0733-00	0.4 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	12/23/2017	

Part 7 of 7

AYPANAL EX

acetaminophen tablet

Product Information

Item Code (Source) NDC:0498-2110

Route of Administration ORAL

Active Ingredient/Active Moiety

0	<i>5</i>			
	Ingredient Name		Basis of Strength	Strength
ACETAMINO PHEN (UNII: 362091)	TL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients

Ingredient Name	Strength	
PO VIDO NE (UNII: FZ989 GH94E)		
MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)		
STARCH, CORN (UNII: O8232NY3SJ)		
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)		
STEARIC ACID (UNII: 4ELV7Z65AP)		

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	12mm
Flavor		Imprint Code	FR1
Contains			

Packaging

	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:0498-2110-01	2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	01/02/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/18/2018	

Establishment			
Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, INC		079287321	pack(0498-4376)

Establishment			
Name	Address	ID/FEI	Business Operations
Ultra Seal Corporation		085752004	manufacture(0498-0114, 0498-2110)

Establishment			
Name	Address	ID/FEI	Business Operations
Water-Jel Technologies		155522589	manufacture(0498-0730, 0498-0903)

Establishment				
Name	Address	ID/FEI	Business Operations	
Changzhou Maokang Medical		421317073	manufacture(0498-0143, 0498-0501)	

Establishment					
Name	Address	ID/FEI	Business Operations		
Safetec of America Inc		874965262	manufacture(0498-0733)		

Revised: 6/2019 Honeywell Safety Products USA, INC